

ABRASA- tetracycline hydrochloride spray
Azimuth Pharma, LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Abrasa Wound Spray

Drug Facts

Active ingredient (in each gram)

Tetracycline hydrochloride 30 mg

Purpose

First aid Antibiotic

Use

- First aid to help prevent skin infection in minor cuts, scrapes, and burns.

Warnings

For external use only

Do not use

- In the eyes or apply over large areas of the body
- Longer than 1 week unless directed by a doctor

Ask a doctor before use if

you have deep or puncture wounds, animal bites, or serious burns.

Stop use and ask doctor if

condition persists or gets worse.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Clean the affected area
- Spray a small amount of this product on the area 1 to 3 time daily
- May be covered with a sterile bandage

Other information

- Stop use if product is misused: if the bottles is left open the liquid will tend to turn dark over time.
- Contains no alcohol, no animal ingredients.
- Blended for typical skin color.
- May stain cloth.

Inactive ingredients

Citric Acid, Dimethyl Sulfoxide, Fulvic Acid, Water, Zinc Oxide

Package Labeling:

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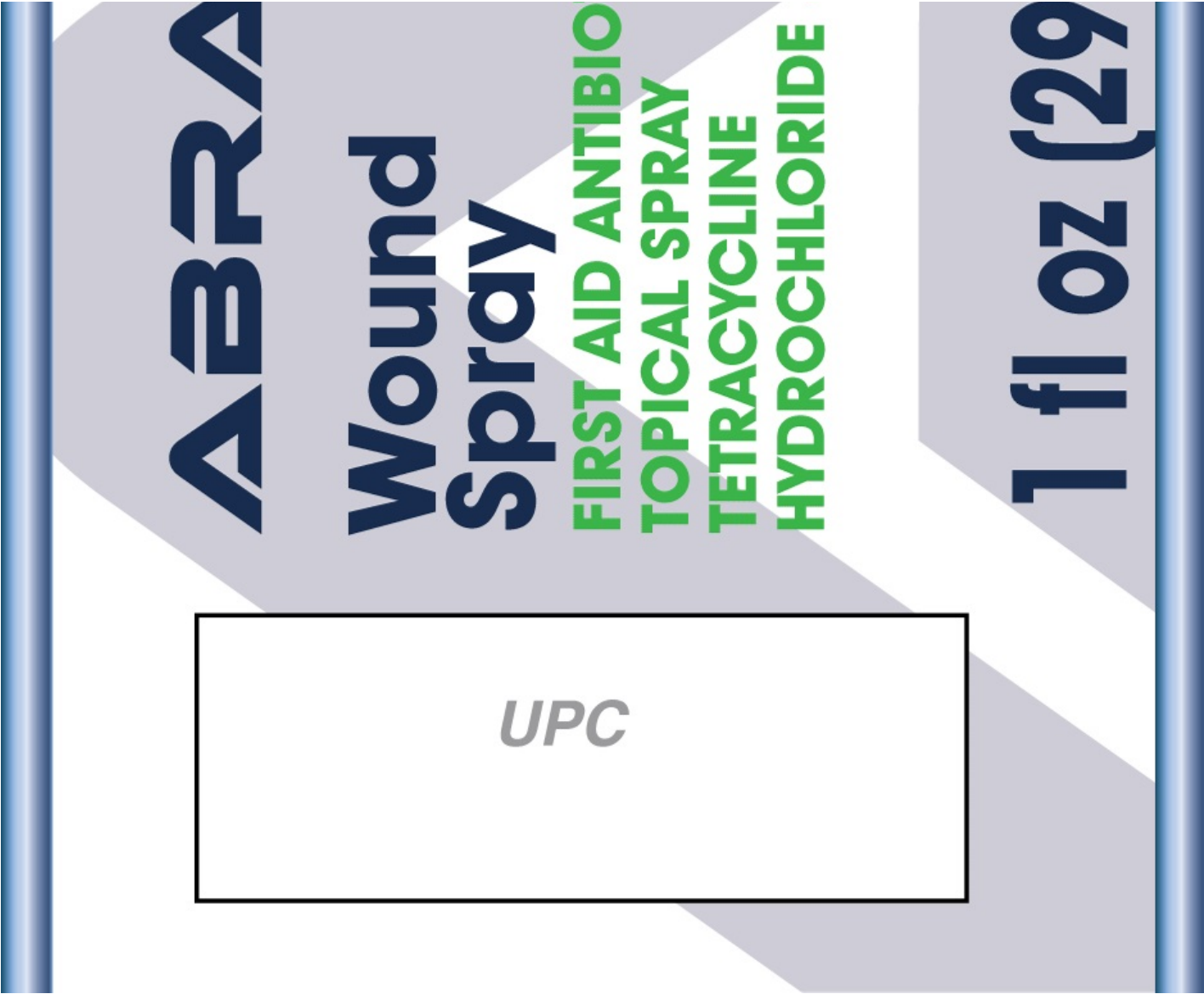
DISTRIBUTED BY: CELLCORE BIOSCIENCES
3750 E. PEWTER FALLS ST., STE 100 MERIDIAN, ID 83642
TO REPORT A SERIOUS ADVERSE EVENT, CONTACT
CELLCORE BIOSCIENCES AT THE ADDRESS ABOVE.
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SA

TIC

30 MG

1.5 mL



ABRASA

tetracycline hydrochloride spray

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81489-001
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
TETRACYCLINE HYDROCHLORIDE (UNII: P6R62377KV) (TETRACYCLINE - UNII:F8VB5M810T)		TETRACYCLINE HYDROCHLORIDE	30 mg in 1 mL
Inactive Ingredients			
Ingredient Name			Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)			

DIMETHYL SULFOXIDE (UNII: YOW8V9698H)	
FULVIC ACID (UNII: XII14C5FXV)	
WATER (UNII: 059QF0KO0R)	
ZINC OXIDE (UNII: SOI2LOH54Z)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81489-001-01	29.5 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/30/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	01/30/2021	

Labeler - Azimuth Pharma, LLC (032996180)

Revised: 4/2021

Azimuth Pharma, LLC